

Salmoporc, lyophilisate for oral suspension for pigs

Authorised

- Salmonella enterica, subsp. enterica, serovar Typhimurium, Live

Product identification

Medicine name:

Salmoporc lyophilisate for oral suspension for pigs

Salmoporc, lyophilisate for oral suspension for pigs

Active substance:

Salmonella enterica, subsp. enterica, serovar Typhimurium, Live

Target species:

Pig

Route of administration:

Oral use

Product details

Active substance and strength:

Salmonella enterica, subsp. enterica, serovar Typhimurium, Live

8.00 log₁₀ colony forming unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral lyophilisate

Withdrawal period by route of administration:**Oral use:**

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Pig

- Meat and offal. 6 week

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AE02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Cardboard box containing one 10 ml glass vial (type I) with rubber stoppers and aluminium crimp caps containing 200 doses (1 dose = 1 ml) of lyophilisate.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Similar biological application (Article 13(4) of Directive No 2001/82/EC)

Marketing authorisation holder:

Ceva Sante Animale

Marketing authorisation date:

18/04/2019

Manufacturing sites for batch release:

Ceva-Phylaxia Veterinary Biologicals Co. Ltd
IDT Biologika GmbH

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10815/064/002

Date of authorisation status change:

18/04/2019

Reference member state:

Netherlands

Procedure number:

NL/V/0247/002

Concerned member states:

Austria Belgium Czechia Denmark Hungary Ireland Italy Portugal Romania
Slovakia United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

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